IN THE CLAIMS:

Please amend the claims as follows:

Claim 1 (Original): An ABC transporter inhibitor which comprises as an active ingredient a cyclic depsipeptide or its optical isomer or racemate of the formula (I):

wherein R¹, R³ and R⁵ are each independently a group selected from linear or branched alkyl having up to 8 carbon atoms; hydroxyalkyl; alkanoyloxyalkyl; alkoxyalkyl; aryloxyalkyl; mercaptoalkyl; alkylthioalkyl; alkylsulfinylalkyl; alkylsulfonylalkyl; carboxyalkyl; alkoxycarbonylalkyl; arylalkoxycarbonylalkyl; carbamoylalkyl; aminoalkyl; alkylaminoalkyl; dialkylaminoalkyl; guanidinoalkyl; alkoxycarbonylaminoalkyl;

9-fluorenylmethoxycarbonyl(Fmoc)aminoalkyl; alkenyl; cycloalkyl; cycloalkylalkyl; and arylalkyl optionally substituted with halogen, hydroxy, alkyl, or alkoxy, and R², R⁴ and R⁶ are each independently a group selected from linear or branched alkyl having up to 8 carbon atoms; hydroxyalkyl; alkanoyloxyalkyl; alkoxyalkyl; aryloxyalkyl; alkylthioalkyl; alkylsulfinylalkyl; alkylsulfonylalkyl; carboxyalkyl; alkoxycarbonylalkyl; arylalkoxycarbonylalkyl; carbamoylalkyl; aminoalkyl; alkylaminoalkyl; dialkylaminoalkyl; alkoxycarbonylaminoalkyl; alkenyl; cycloalkyl; cycloalkylalkyl; and aryl or arylalkyl which are optionally substituted with halogen, hydroxy, alkyl, or alkoxy.

Claim 2 (Original): The ABC transporter inhibitor according to claim 1, wherein the cyclic depsipeptide is a compound of the formula (II):

wherein R¹, R³ and R⁵ are each independently linear or branched lower(C₁₋₄)alkyl.

Claim 3 (Original): The ABC transporter inhibitor according to claim 2, wherein the groups represented by R1', R3' and R5' are linear or branched propyl or butyl.

Claim 4 (Original): The ABC transporter inhibitor according to claim 3, wherein R¹ and R³ are each isopyropyl, and R⁵ is any one of the groups selected from isopropyl, sec-butyl, and isobutyl.

Claim 5 (Currently Amended): The ABC transporter inhibitor according to any one of claims 1 to 4 claim 1, wherein the ABC transporter is MDR protein.

Claim 6 (Currently Amended): The ABC transporter inhibitor according to any one of claims 1 to 4 claim 1, wherein the ABC transporter is CDR1 or CDR2 protein of Candida yeast.

Claim 7 (Currently Amended): The ABC transporter inhibitor according to any one of elaims 1 to 4 claim 1, wherein the ABC transporter is PDR5 protein of Saccharomyces yeast.

Claim 8 (Original): An inhibitor against the acquisition of drug resistance, which comprises as an active ingredient a cyclic depsipeptide or its optical isomer or racemate of the formula (I):

wherein R¹, R³ and R⁵ are each independently a group selected from linear or branched alkyl having up to 8 carbon atoms; hydroxyalkyl; alkanoyloxyalkyl; alkoxyalkyl; aryloxyalkyl; mercaptoalkyl; alkylthioalkyl; alkylsulfinylalkyl; alkylsulfonylalkyl; carboxyalkyl; alkoxycarbonylalkyl; arylalkoxycarbonylalkyl; carbamoylalkyl; aminoalkyl; alkylaminoalkyl; dialkylaminoalkyl; guanidinoalkyl; alkoxycarbonylaminoalkyl;

9-fluorenylmethoxycarbonyl(Fmoc)aminoalkyl; alkenyl; cycloalkyl; cycloalkylalkyl; and arylalkyl optionally substituted with halogen, hydroxy, alkyl, or alkoxy, and R², R⁴ and R⁶ are each independently a group selected from linear or branched alkyl having up to 8 carbon atoms; hydroxyalkyl; alkanoyloxyalkyl; alkoxyalkyl, aryloxyalkyl; alkylthioalkyl; alkylsulfinylalkyl;

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 $alkyl sulfonylalkyl;\ carboxyalkyl;\ alkoxycarbonylalkyl;\ arylalkoxycarbonylalkyl;$

carbamoylalkyl; aminoalkyl; alkylaminoalkyl; dialkylaminoalkyl; alkoxycarbonylaminoalkyl;

alkenyl; cycloalkyl; cycloalkylalkyl; and aryl or arylalkyl which are optionally substituted with

halogen, hydroxy, alkyl, or alkoxy.